Effect of Physical Training on the Adipose Tissue of Diet-induced Obesity Mice: Interaction Between Reactive Oxygen Species and Lipolysis

Abstract

It is well known that high-fat diets (HFDs) induce obesity and result in an increase in oxidative stress in adipose tissue, which leads to an impairment of fat mobilization by a downregulation of the lipases, such as hormone-sensitive lipase (HSL) and adipose triglyceride lipase (ATGL). On the other hand, exercise training leads to a reduction in adipose tissue and an improvement of antioxidant status and the lipolytic pathway. Our aim was to examine the influence of exercise and moderate intensity training on oxidative stress parameters and the relationship between the proteins involved in the lipolysis of animals subjected to a high-fat fed diet. Twenty-four mice were used and divided into 4 groups (n = 6): standard diet (SD); standard diet plus exercise (SD + Ex); high-fat diet (HFD); and high-fat diet plus exercise (HFD + Ex). The animals received HFD for 90 days and submitted to a daily training protocol in swinging. The animals were euthanized 48 h after the last session of exercise. White adipose tissue epididymal fat was excised on fatty acid (FA) supply, FA esterification to triglycerides (TG), and TG breakdown, or lipolysis. Both adipose triglyceride lipase (ATGL) and hormone-sensitive lipase (HSL) have the capacity for TG degradation by cleaving the ester bond, governing the lipolysis pathway in adipose tissue [6]. Adipose tissue lipolysis has received much attention over the past 10 years because of its altered regulation in obesity.

Introduction

Obesity is a public health problem that affects individuals of all social classes, gender, and age. Genetic predisposition, several environmental and behavioral factors are associated with the pathophysiology of obesity, and are recognized as risk factors for lifestyle-related diseases, such as cardiovascular disease and diabetes [1]. On the other hand, exercise training has been used as an important therapeutic resource for both the treatment and prevention of several diseases. It is now known that exercise training results in less body weight, adipose tissue depots via the regulation of lipases, and induces anti-inflammatory and antioxidant profiles [2–5]. Adipose tissue fat stores are mainly dependent on fatty acid (FA) supply, FA esterification to triglycerides (TG), and TG breakdown, or lipolysis. Both adipose triglyceride lipase (ATGL) and hormone-sensitive lipase (HSL) have the capacity for TG degradation by cleaving the ester bond, governing the lipolysis pathway in adipose tissue [6]. Adipose tissue lipolysis has received much attention over the past 10 years because of its altered regulation in obesity.
homeostasis. However, ROS levels can increase dramatically due to several stimuli, and this may result in significant damage to cell structures [8]. Several studies show the involvement of ROS in the pathophysiology of obesity [2,9–11] and suggest that excessive ROS formation results in oxidative stress and inflammation induces significant changes in the gene expression of adipocytes, lipid metabolism, and the content of adipocytes. More recently, Krawczyk et al. [12] have postulated the contribution of oxidative stress to the control lipolysis in the adipose tissue. However, this interaction was never examined in obese animals that were concomitantly submitted to physical training protocol. Here, we examined the molecular and biochemical mechanisms involved in obesity, investigated the oxidative stress parameters and proteins involved in lipolysis in adipose tissue, and explored the effects of exercise training as nonpharmacologic therapy.

Materials and Methods

Samples
Twenty-four 2-month-old male Swiss mice from the colony maintained by the Universidade do Extremo Sul Catarinense (UNESC), Criciúma, Santa Catarina, Brazil, were used in this study. The animals were randomized into 4 groups (n=6): standard diet (SD); standard diet plus exercise (SD + Ex); high-fat diet (HFD); and high-fat diet plus exercise (HFD + Ex). They were housed 4 per cage in an animal room under a 12-h light-dark cycle at 22 ± 1°C and received a chow diet and water ad libitum. The experiments were carried out after a one-week acclimation period. All procedures were conducted in accordance with regulation no. 11794/08 (DOU 196, Section 1, October 2008) and approved by the local ethics committee.

Diet and exercise protocol
Composition of the experimental diet was according to Noeman and colleagues [11]. The diet applied was purchased from Nuvital Nutrientes SA, Brazil (Table 1). All animals had free access to water and food for 90 days. Thirty days after having started the high-fat diet, the animals were submitted to a daily training protocol. The exercise groups performed a continuous swimming program 5 days/week for 8 weeks. The mice swam for two 30-min sessions separated by a 5-min break. The untrained animals were placed in an empty swimming pool for the same amount of time as the exercise group for the entire duration of the 8 weeks. After 4 weeks, a tail weight (5% of body weight) was used to increase the intensity of training. Training was conducted in a 120 × 60 × 50 cm pool with 10 × 15 × 50 cm lanes and containing 37.5 cm³ water that was temperature controlled at 30–32°C.

Weight and exercise control
Body weight was measured both before and immediately after the last training sessions using a precision semi-analytic balance (Gehaka model BK300, precision 0.01 g). Training intensity was controlled by the blood lactate level. Before and immediately after the first, twentieth, and last training sessions, approximately 25 μl blood samples were collected from the caudal vein of animals for lactate measurement using a lactometer (Accusport). Control animals were submitted to one training session only, for lactate measurement sampling. Body weight was measured during all experimental periods using a precision semi-analytic balance (Gehaka model BK300, precision 0.01 g).

Euthanasia
Forty-eight hours after the last exposure session, mice were killed by cervical dislocation, and adipose tissue of visceral regions (epididymal fat) was surgically excised. Next, an aliquot of these tissues was homogenized in a specific buffer and used for biochemical analyses and intracellular proteins by Western blotting. The remaining material was stored in a freezer at ~80°C for further analysis.

Homogenization of sample for Western blotting
The samples were homogenized in a buffer containing 1% Triton X 100, Tris 100 mM (pH 7.4), sodium pyrophosphate 100 mM, EDTA 100 mM, sodium vanadate 10 mM, PMFS 2 mM, and aprotonine 0.1 mg/ml at 4°C. The homogenate was then centrifuged at 11000 rpm for 40 min. The supernatant was used to determine the concentration of protein, and later, total extract was determined in the same material, using a specific antibody.

Western blotting
Aliquots containing 250 μg of protein (per sample) were applied on a 1.5 mm thick polyacrylamide gel. Electrophoresis was conducted in a minigel cell (Bio Rad, Mini-Protean) with an electrophoresis buffer that had been diluted earlier. SDS-PAGE was initially conducted at 25 V (on the stacking gel) and 120 V (until the end of the resolution gel). Next, proteins separated by SDS-PAGE were transferred to a nitrocellulose membrane using an electrotransference minigel device (Bio Rad), and the running buffer solution was maintained at 120 V for 2 h under continuous refrigeration using ice. Nitrocellulose membranes containing run proteins were incubated in a blocking solution for 2 h at room temperature to reduce the nonspecific protein ligation. Then, the membranes were washed thrice (for 10 min each time) in wash buffer and incubated in specific antibodies (anti-superoxide dismutase (SOD), anti-catalase (CAT), anti-ATGL (adipose triglyceride lipase), anti-CGI-58 (comparative gene identification 58), and anti-FAS (fatty acid synthase antibodies)) under constant shaking overnight at 4°C. Next, the membranes were washed again thrice (for 10 min each time) in wash buffer and incubated in secondary antibodies conjugated with peroxide for 2 h at room temperature. Excess secondary antibody was washed with wash buffer, and then, the membranes were incubated in enzyme substrate for 2 min and exposed to an X-ray film (Kodak XAR, Rochester, NY, USA) with an intensifier (Cronex Lightning...
Absorbance was read at 750 nm [17].

Protein content was assayed using bovine serum albumin as a standard. A folin phenol reagent (phosphomolybdic-phosphovanadyl reagent) was added to bind the protein. The bound reagent was determined by spectrophotometry at 370 nm with a coefficient of 22 000 molar$^{-1}$ and was expressed as nmol/mg protein [16].

### Oxidative damage

Thiobarbituric acid reactive species (TBARS) was measured as a marker of lipid peroxidation by spectrophotometry at 532 nm. Results were expressed as nmol TBARS/mg protein [15], and the generation of carbonyl groups in the reaction with 2,4-dinitrophenylhydrazine was used as a marker of the oxidation of proteins. Carbonyl content was determined by spectrophotometry at 370 nm with a coefficient of 22 000 molar$^{-1}$ and was expressed as nmol/mg protein [16].

### Antioxidant enzyme activities

Superoxide dismutase (SOD) activity was determined according to the method described by Bannister and Calabrese [13]. The enzymatic activity estimation occurs by adrenaline autooxidation inhibition read at 480 nm in a spectrophotometer. Enzyme activity was expressed as U/mg protein. To determine catalase (CAT) activity, the sample was sonicated in a 50 mM phosphate buffer, and the resulting suspension was centrifuged at 3000 g for 10 min. The supernatant was used for enzyme assay. CAT activity was measured using the rate of decrease in hydrogen peroxide absorbance at 240 nm, determined according to the method described in [14]. Enzyme activity was expressed as U/mg protein.

### Oxidative damage

Protein content was assayed using bovine serum albumin as a standard. A folin phenol reagent (phosphomolybdic-phosphotungstic reagent) was added to bind the protein. The bound reagent was slowly reduced and changed from yellow to blue. Absorbance was read at 750 nm [17].

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### Results

#### Body weight and training control

Table 2 shows body weight after exposure to a high-fat diet and exercise training protocol. Body weight in the HFD showed increased 37 % (p < 0.001), whereas HFD + Ex exhibited increased 15 % (p < 0.05) when compared with SD. In addition, HFD + Ex exhibits a body weight of 19% minor (p < 0.05) in relation to HFD. Lactate level increased in both groups treated with a standard diet and a high-fat diet, in relation to basal values.

### Antioxidant enzymes

Fig. 1 shows that SOD activity and level changed after exercise training with regard to animals' exposure to HFD. The SOD level in adipose tissue showed an increase in HFD (p < 0.05) compared to SD, whereas the HFD + Ex group showed an increase in SOD activity relative to SD, HFD, and SD + Ex (p < 0.05). In relation to CAT (Fig. 2), the results showed an increase in activity in the HFD + Ex in relation to HFD (p < 0.05), whereas the protein level of enzymes in adipose tissue was increased in the HFD + Ex group compared to SD, HFD, and SD + Ex (p < 0.05).

### Oxidative damage

Fig. 3 shows the values of TBARS and carbonyl protein. In the adipose tissue, the lipid peroxidation and protein oxidation were significantly increased in the HFD (p < 0.01) compared to the SD group. In contrast, exercise training leads to a decrease in both parameters (p < 0.01).

#### Table 2 Body weight and training control.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Body weight (mg)</th>
<th>Blood lactate (nmol/l)</th>
<th>Mean ± SEM</th>
<th>Mean ± SEM</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Basal</td>
<td>1st session</td>
<td>20th session</td>
</tr>
<tr>
<td>SD</td>
<td>40.43 ± 1.08</td>
<td>2.13 ± 0.19</td>
<td>3.92 ± 0.13*</td>
<td>4.27 ± 0.25*</td>
</tr>
<tr>
<td>HFD</td>
<td>55.29 ± 2.00*</td>
<td>4.68 ± 1.83*</td>
<td>4.10 ± 0.10*</td>
<td>4.60 ± 0.20*</td>
</tr>
<tr>
<td>SD + Ex</td>
<td>37.14 ± 0.90</td>
<td>3.92 ± 0.13*</td>
<td>4.27 ± 0.25*</td>
<td>4.13 ± 0.27*</td>
</tr>
<tr>
<td>HFD + Ex</td>
<td>46.88 ± 1.83*</td>
<td>4.27 ± 0.25*</td>
<td>4.13 ± 0.27*</td>
<td></td>
</tr>
</tbody>
</table>

Blood lactate level (nmol/l) of mice exposed to physical training, a high-fat diet (HFD), and a standard diet (SD). Values are presented as mean ± SEM, and p < 0.05 was considered as the minimum significance between the groups.

* Different from a standard diet (SD). ▲ Different from a high-fat diet (HFD). ▼ Different from a standard diet plus exercise (SD + Ex)

#### Fig. 1 Activity and level of superoxide dismutase in epididymal fat (a and b) of mice submitted to training after exposure to a high-fat diet. Values are presented as mean ± SEM, and p < 0.05 was considered as the minimum significance between the groups. ▲ Different from a standard diet (SD), ▲ different from a high-fat diet (HFD), and ▼ different from a standard diet plus exercise (SD + Ex).
Lipolytic and lipogenic proteins

**Fig. 4** shows the data of ATGL, CGI-58, and FAS, respectively. ATGL protein levels were increased in the HFD + Ex (p < 0.05) compared to all the groups. CGI-58 protein levels were reduced in all the groups (p < 0.05) in relation to the SD groups. FAS protein levels increased in the HFD (p < 0.05) relative to the SD and HFD + Ex groups, whereas they exhibited decreases in the SD + Ex and HFD + Ex (p < 0.05) than in the SD group.

**Discussion**

Our results indicate that obesity induces great alterations with regard to several biochemical and molecular parameters in adipose tissue, whereas physical training rebalances these alterations, improving antioxidant profile, and increasing ATGL protein levels, which induces minor weight gain. Here, we have demonstrated that, at least in part, changes in oxidative stress are associated with the lipolysis process in adipose tissue in high-fat fed mice.
Lactate content was used to monitor changes in the physical capacity of animals and to control the training intensity. Changes in muscle mass alter the energetic metabolism due to differences in oxidative and glycolytic capacities for adenosine triphosphate (ATP) turnover. Thus, this marker may be considered as an indicator of the intensity of physical effort during training. Our results show an increase in lactate content in all groups after training sessions (immediately after the first, the twentieth, and the last training session). This increase suggests that the protocol utilized reflects aerobic training and can result in an impact on oxidative metabolism.

Obesity is closely associated with ROS, and studies have shown that the unbalance between oxidant production and the antioxidant defense system is critically involved in the pathogenesis of obesity in the human [9, 10] and experimental model [11, 18]. It has been reported that obesity may induce systemic oxidative stress, and oxidative stress is associated with the overproduction of proinflammatory adipokines [19]. In contrast, antioxidant defense markers are lower depending on the amount of body fat [20, 21]. Our data corroborated with those found in the literature, which showed that adipose tissue in high-fat fed diet mice exhibits an increase in ROS concentrations [22]. The increase in obesity-associated ROS is probably due to the relationship between obesity and chronic inflammation, because adipocytes and preadipocytes have been identified as a source of proinflammatory cytokines, including TNF-α, IL-1β, and IL-6 [23, 24], and these cytokines serve as potent stimulators for the production of ROS by macrophages and monocytes [25]. In addition, it is observed that disruption in the lipolysis pathway leads to accumulated TAG in the adipose tissue, which favors an increase in this tissue and aggravates obesity. Adipocyte triglyceride lipase (ATGL) and hormone-sensitive lipase (HSL) assume triglyceride hydrolysis activities in the adipose tissue. Lipolysis, which is coordinated by lipases, occurs through catecholamine stimulation of β-adrenergic receptors and the subsequent activation of protein kinase A (PKA) [26]. In mice, the inhibition of HSL and ATGL has been shown to promote diet-induced obesity and its associated metabolic defects [27]. In humans, both HSL and ATGL protein levels are decreased in the adipose tissue from obese patients compared with that from lean individuals [28, 29].

Our data show that the FAS protein level is increased in the adipose tissue in high-fat-fed diet mice, which favors the synthesis of fatty acids, whereas there are no changes observed in the ATGL protein level. The limitation of the present study was that phosphorylation of the enzyme was not examined. Taken together, our data suggest that an increase in oxidative stress is accompanied by the disruption lipolysis process in the adipose tissue high-fat fed diet mice. On the other hand, exercise training has been associated with an improvement in the antioxidant defense system and a decrease in oxidative damage in the brain [30], lung [31], liver [32], muscle [33], kidney [34], heart [35], adipose tissue [2], and other tissues as well as the reduction in obesity [36, 37].

In the present study, we demonstrated that exercise training prevents marked weight adipose tissue gain, oxidative damage, and recovery of the ATGL protein level in adipose tissue. These data suggest that the protector effects of exercise training have a link between reduced ROS and the lipolysis pathway. This interaction has been recently postulated by Krawczyk et al. [12], who demonstrated that ROS facilitates the translocation of HSL to the lipid droplet during lipolysis in human adipocytes.
fat-fed rats and might, at a first glance, simply reflect a reduction in fat mass in the adipose tissue. In contrast to HSL, ATGL appears to reside on the lipid droplet surface independent of PKA activation [43]. Rather than its activation being controlled by phosphorylation and translocation, ATGL is activated at least 20-fold by interactions with CGI-58, a member of an esterase/lipase family of proteins [44]. Comparative Gene Identification-58 (CGI-58) is also known as α/β-hydrolase domain-containing protein 5 (Abhd5). CGI-58 lacks lipase activity itself, but activates the lipase activity of ATGL likely via protein–protein interaction. Fluorescence resonance energy transfer studies support a model in which CGI-58 binds to perilipin in adipocytes under basal conditions but is released from perilipin on lipolytic stimulation [45]. We suggest that, despite reduced CGI-58 protein levels in the HFD, the interactions between CGI-58 and perilipin have been increased, but cannot affirm this situation. More studies are needed to confirm the suggestion. In summary, the results of this study suggest that antioxidant enzymes play a significant role in the regulation of obesity, but are not sufficient to reduce the oxidative damage induced by exposure to HFD. However, since we did not determine anti-inflammatory cytokines and NAPH oxidase, the exact role of physical training in the modulation of oxidative stress in animals exposed to HFD should be confirmed in future studies.

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Conflict of Interest
The authors have no conflict of interest.

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